

SICKLE-CELL ANEMIA



erythrocytes
of a healthy person



human erythrocytes
of a patient with sickle-cell anemia

Sickle Cell Anemia

Elena Ledger

A red circular graphic with a gradient, appearing as a semi-circle or a partial circle, located to the right of the light blue bar.

Sickle Cell Anemia:

Sickle Cell Anemia Judy Monroe Peterson,2008-08-15 Sickle cell anemia is a genetic disease of the blood It is caused by a defect in one gene of a person Genes are the elements in cells that carry the information that determines traits such as hair or eye color In sickle cell anemia a defect in the gene controls how hemoglobin is made This defect can be passed from parents to their children Students explore the history of sickle cell anemia the pioneering doctors who studied its cause and early treatments They also investigate hemoglobin S who gets sickle cell and how the gene mutation is inherited They learn about the different types of sickle cell disease and treatments including blood transfusions and bone marrow transplants and some of the promising new research in medicines and gene therapy Sickle Cell Anemia Jane S. Lin-Fu,1975 **Sickle Cell Anemia** Jane S. Lin-Fu,1975 **Fact Sheet, Sickle Cell Anemia** National Sickle Cell Disease Program,National Heart and Lung Institute,1975 *Sickle Cell Anemia: From Basic Science to Clinical Practice* Elena Ledger,2021-11-16 A sickle cell disease is a group of blood disorders that a person inherits from parents They usually occur when the person inherits two abnormal copies of the hemoglobin gene Sickle cell anemia is the most common type of sickle cell disease It causes an abnormality in hemoglobin the oxygen carrying protein found in red blood cells The problems due to sickle cell anemia begin to appear around 5 to 6 months of age It leads to problems such as sickle cell crisis swelling in hands and feet stroke and bacterial infections The care of people suffering from sickle cell anemia includes infection prevention with vaccination and antibiotics folic acid supplementation and pain medication A bone marrow transplant is also used in certain cases This book consists of contributions made by international experts It contains some path breaking studies in sickle cell anemia It will serve as a valuable source of reference for graduate and post graduate students **Sickle Cell Anemia** ,1990

Understanding Sickle Cell Disease Miriam Bloom,1995 Explains how sickle cell anemia is inherited describes its symptoms and treatment and discusses the search for a cure *What You Can Do About Sickle Cell Disease* Monique Vescia,Alvin Silverstein,Virginia Silverstein,2015-07-15 What is sickle cell disease What damage can it cause Can anyone get it Can it be treated or prevented This text explains a rather complex and potentially deadly condition through case studies and interviews and presents the most current research and treatments Sickle Cell Anemia Alvin Silverstein,Virginia B. Silverstein,Laura Silverstein Nunn,1997 A hereditary disease that is suffered by more than 60 000 Americans today sickle cell anemia affects red blood cells Includes the history of the disease case studies and treatments available today Sickle Cell Anemia Ruth Bjorklund,1997-01-01 Provides comprehensive information on the causes treatment and history of sickle cell anemia *I Saw His Face Before Me* Samuel Burns,Patricia Burns,2013-04-05 I Saw His Face Before Me is the story of the late Heather Anese Burns her surviving parents Samuel and Patricia Burns and her much loved brother Keenan This achingly honest moving account of two deserved lives renders page upon page of wisdom knowledge and compassion compelling us to face the challenges and celebrate the successes of Heather s and Keenan s journey with Sickle Cell Anemia

Encircled in a heartwarming message the authors offer riveting eye opening insights into parenting children and adults with incurable diseases knowing each first time experience might be a first time once I can do all things through Christ which strengthens me Philippians 4:13 Heather's passion for all things Godly and her love for education and learning inspired her family and others who loved her to establish the Heather Burns Memorial Scholarship Fund Learn more about the HBMSF at www.hbmsf.org

Planning for Sickle Cell Anemia in Pennsylvania Pennsylvania. Irvis Committee on Sickle Cell Anemia, 1973 *Sickle Cell Anemia: New Insights for the Healthcare Professional: 2011 Edition*, 2012-01-09 Sickle Cell Anemia New Insights for the Healthcare Professional 2011 Edition is a ScholarlyBrief that delivers timely authoritative comprehensive and specialized information about Sickle Cell Anemia in a concise format The editors have built Sickle Cell Anemia New Insights for the Healthcare Professional 2011 Edition on the vast information databases of ScholarlyNews You can expect the information about Sickle Cell Anemia in this eBook to be deeper than what you can access anywhere else as well as consistently reliable authoritative informed and relevant The content of Sickle Cell Anemia New Insights for the Healthcare Professional 2011 Edition has been produced by the world's leading scientists engineers analysts research institutions and companies All of the content is from peer reviewed sources and all of it is written assembled and edited by the editors at ScholarlyEditions and available exclusively from us You now have a source you can cite with authority confidence and credibility More information is available at <http://www.ScholarlyEditions.com>

Handling Sickle Cell Disease Yvette LaPierre, 2021-08-01 This book explores sickle cell disease and how people with the disease manage it It examines how sickle cell affects daily life work and school and it explains the latest treatments available Features include a glossary web resources source notes and an index Aligned to Common Core Standards and correlated to state standards Essential Library is an imprint of Abdo Publishing a division of ABDO [Sickle Cell Anemia](#) George Beshore, 1994-01-01 Looks at the history and symptoms of sickle cell anemia describes how it is treated and discusses current research

Directory of National, Federal, and Local Sickle Cell Disease Programs National Heart, Lung, and Blood Institute. Sickle Cell Disease Branch, 1978 Sickle cell facilities in the United States the Bahamas and Puerto Rico Entries arranged under sections titled Comprehensive sickle cell centers Sickle cell screening and education clinics National centers for family planning services Veterans administration hospitals Job Corps regions and Public and private organizations Each entry gives name address and telephone number [Sickle Cell Anemia: the Neglected Disease; Community Approaches to Combating Sickle Cell Anemia](#) "San Francisco Symposium "Sickle Cell Anemia - the neglected disease, 1973 *Hope and Destiny* Allan F. Platt, Alan Sacerdote, 2006 For anyone who has or is predisposed to sickle cell disease this informative and compassionate guide provides all the facts patients loved ones and caregivers need to know in order to reduce symptoms relieve pain and help patients and their support circle better understand the cause and growth of the disease Divided into different sections to address the changing complications posed by the disease at each stage of life this book emphasizes the need for offering

emotional and spiritual consolation to those who suffer from sickle cell disease or witness the suffering of a love one Topics include the complex causes of sickle cell disease the most current treatment options facts on genetic counseling pain assessment and management resources and strategies to lower the likelihood of pain crises *Sickle Cell Disease: Concerns and Challenges* Sabrina Kelley,2020-09-22 Sickle cell disease SCD is a group of inherited blood disorders Sickle cell anemia SCA is the most common form of SCD It is characterized by an abnormality in hemoglobin found in red blood cells This results in rigid sickle shaped red blood cells SCA typically develops symptoms like anemia bacterial infections stroke pain swelling in the hands and feet etc The gene defect associated with sickle cell disease is a single nucleotide mutation Normal red blood cells are elastic thereby allowing cells to deform and pass through capillaries In SCD low oxygen tension promotes sickling of red blood cells which fail to return to the normal shape when the ideal oxygen tension is restored These cells are therefore not able to pass through narrow capillaries leading to ischemia and vessel occlusion SCA can lead to complications such as stroke severe bacterial infections cholelithiasis avascular necrosis osteomyelitis acute papillary necrosis etc This book is compiled in such a manner that it will provide in depth knowledge about the concerns and challenges in the management of sickle cell disease It brings forth some of the most innovative concepts and elucidates the unexplored aspects of this disease It is meant for students who are looking for an elaborate reference text on sickle cell disease **The Sickle Cell Anemia Update** Alvin Silverstein,Virginia B. Silverstein,2006 Sickle Cell Anemia is discussed with regard to causes detection and treatment and prevention

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